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# IN THE CLAIMS ATTACHED TO THE INTERNATIONAL PRELIMINARY EXAMINATION REPORT:

Please rewrite claims 2, 4-8, 10, 11 and 13-37 as shown below in the detailed listing of all claims which are, or were, in this application:

- 1. (Original) A method for preparing a sol-gel derived SiO<sub>2</sub> monolith, preferably with a minimum diameter of ≥ 0.5 mm, coating, preferably with a thickness of < 0.5 mm, or particle, preferably with a maximum diameter of ≤ 100 μm, with a very fast bioresorption rate, said SiO<sub>2</sub> optionally comprising a specific percentage or percentages of a biologically active agent or agents other than the SiO<sub>2</sub> itself with or without protective agent or agents for said biologically active agent or agents, wherein method a sol-gel derived SiO<sub>2</sub> is prepared from a sol comprising water, an alkoxide or inorganic silicate and a lower alcohol, i.e. an alcohol with ≤ 4 carbons, using a mineral acid or a base as a catalyst, preferably a mineral acid, and said sol is aged and dried **characterised** in that
- a) in the sol the starting
  - i) pH is from 0.05 to 2.5, preferably 1.5 to 2.5, most preferably 2.0,

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- ii) molar ratio of water to the alkoxide or inorganic silicate is 0.5 to 2.5; preferably 1.5 to 2.5,
- iii) molar ratio of alcohol to the alkoxide or inorganic silicate is  $\geq$  0.5, preferably  $\geq$  1.0; and
- b) either,
  - i) the sol is, without induced changes of sol composition,
    - let to gel spontaneously at a temperature of ≤ 25

      °C or an elevated temperature of 65 °C to 90 °C,

      preferably at an elevated temperature of 65 °C to

      90 °C, or
    - gelation of the sol is done by forced drying of the sol, or
  - ii) a change or changes of sol composition are induced after sol ageing but before gel formation, said change or changes of sol composition optionally comprising addition of said biologically active agent or agents with or without said protective agent or agents, and

the ratio  $t/t_{gel}$  is  $\geq$  0.005, preferably  $\geq$  0.1, most preferably  $\geq$  0.9, wherein

t is the ageing time of the sol, i.e. time from preparation of said sol to the induced changes, and

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 $t_{gel}$  is the time point where the sol would have turned to a gel without the induced changes; and forced drying of the sol is carried out or initiated within a time of  $\leq$  30 minutes, preferably  $\leq$  15 minutes, most preferably  $\leq$  5 minutes, from said induced change or changes.

- 2. (Currently amended) A method for adjusting the bioresorption rate of sol-gel derived  $SiO_2$  monolith, preferably with a minimum diameter of  $\geq$  0.5 mm, coating, preferably with a thickness of < 0.5 mm, or particle, preferably with a maximum diameter of  $\leq$  100  $\mu$ m, optionally comprising a specific percentage or percentages of a biologically active agent or agents other than the  $SiO_2$  itself with or without protective agent or agents for said biologically active agent or agents, characterised in that
- A) a  $SiO_2$  with a very fast bioresorption rate is obtained according to the method of preparing a  $SiO_2$  of claim 1; and
- B) a SiO<sub>2</sub> with a slower bioresorption rate than the very fast bioresorption rate is obtained by correlating a desired biodegradability of a SiO<sub>2</sub> with changes a), b) and/or c) to the said method of preparing a SiO<sub>2</sub> according to claim 1, wherein

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- a) comprises deviating in the sol any of the starting values:
  - i) pH,
  - ii) molar ratio of water to the alkoxide or inorganic silicate, and/or
  - iii) molar ratio of alcohol to the alkoxide or inorganic silicate;

from the values defined in a) i) - iii) of claim 1 said method of preparing a  $SiO_2$ ;

- b) comprises carrying out induced changes by addition of a component or components, including optional addition of the biologically active agent or agents with or without said protective agent or agents, said changes affecting any of the values i) iii) of a) of claim 1 said method of preparing a  $SiO_2$  or a) above if applied by
  - i) not carrying out forced drying, or
  - ii) carrying out or initiating forced drying of the sol later than defined in b) ii) of claim 1 said method of preparing a  $SiO_2$ ; and

- c) comprises deviating the temperature for letting the sol gel spontaneously from the values defined in b) i) of  $\frac{1}{1}$  said method of preparing a  $\frac{1}{1}$  and
- a method for preparing the  $SiO_2$  with said changes to the method correlating with the desired biodegradability is carried out for obtaining the  $SiO_2$  with the desired slower biodegradability.
- 3. (Original) The method according to claim 2 **characterised** in that an alkoxide, preferably tetraethoxysilane (TEOS), is used for preparing the sol-gel derived  $SiO_2$ .
- 4. (Currently amended) The method according to claim 2 or 3 characterised in that that an inorganic silicate, preferably sodium or potassium silicate, is used for preparing the sol-gel derived  $SiO_2$ .
- 5. (Currently amended) The method according to any of claims 2 to 4 claim 2 characterised in that the lower alcohol is ethanol.

- 6. (Currently amended) The method according to any of claims 2 to 5 claim 2 characterised in that the induced change is selected from the group consisting of adding water, adding the alkoxide or inorganic silicate, adding the alcohol, adjusting pH by adding an acid or base, preferably the acid or base used as the catalyst, adding the optional bioactive agent or agents with or without protective agent or agents for said biologically active agent or agents affecting any of the values i) iii) of a) in claim 1 of said method of preparing a SiO<sub>2</sub> or a) of claim 2 said method of adjusting the bioresorption of sol-gel derived SiO<sub>2</sub> if applied, and any combination thereof.
- 7. (Currently amended) The method according to any of claims 2 to 6 claim 2 characterised in that drying of the sol is drying by ambient heat, vacuum drying, electromagnetic drying, acoustic drying, spray-drying or freeze-drying, preferably spray-drying or freeze-drying.
- 8. (Currently amended) The method according to any of claims 2 to 7 claim 2 characterised in that forced drying of the sol is carried out, preferably by spray-drying or freeze-drying.

- 9. (Original) The method according to claim 8 characterised in that forced drying is freeze-drying initiated by freezing the sol.
- 10. (Currently amended) The method according to claim 8 or 9 characterised in that the temperature of the sol is  $\leq$  +90 °C, preferably  $\leq$  +50 °C, most preferably  $\leq$  +40 °C.
- 11. (Currently amended) The method according to any of claims 2 to

  10 claim 2 characterised in that the gel is dried.
- 12. (Original) The method according to claim 11 characterised in that drying of the gel is drying by ambient heat, vacuum drying, electromagnetic drying, acoustic drying, spray-drying or freezedrying, preferably ambient heat or freeze-drying.
- 13. (Currently amended) The method according to claim 11 or 12 characterised in that the gel is dried at a temperature of  $\leq$  700 °C, preferably  $\leq$  50 °C, and most preferably  $\leq$  40 °C.
- 14. (Currently amended) The method according to any of claims 2 to

  13 claim 2 characterised in that a value to be deviated to obtain

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a slower bioresorption rate is the ratio of water to the alkoxide or inorganic silicate, and the more the ratio of water to alkoxide or inorganic silicate is deviated to be higher or lower the slower the bioresorption rate obtained.

- 15. (Currently amended) The method according to any of claims 2 to 14 claim 2 characterised in that a value to be deviated to obtain a slower bioresorption rate is the ratio of alcohol to the alkoxide or inorganic silicate, and the more the ratio is deviated to be higher or lower the slower the bioresorption rate obtained.
- 16. (Currently amended) The method according to any of claims 2 to 15 claim 2 characterised in that a value to be deviated to obtain a slower bioresorption rate is the pH, and the more the pH is deviated to be higher or lower the slower the bioresorption rate obtained.
- 17. (Currently amended) The method according to any of claims 2 to 16 claim 2 characterised in that a biologically active agent or agents is added to the sol before gel formation.

- 18. (Currently amended) The method according to any of claims 2 to  $\frac{17 \text{ claim 2}}{17 \text{ claim 2}}$  characterised in that any of the values pH, molar ratio of water to the alkoxide or inorganic silicate, and/or molar ratio of alcohol to the alkoxide or inorganic silicate is changed to deviate from the ranges defined in claim 1 in said method of preparing a  $\frac{100}{100}$ , a) ic) iii), after sol ageing but before gel formation and/or optional addition of said biologically active agent or agents, and within  $\frac{100}{100}$  minutes, preferably  $\frac{100}{100}$  15 minutes and most preferably  $\frac{100}{100}$  5 minutes from the change forced drying of the sol is carried out or initiated.
- 19. (Currently amended) The method according to any of claims 2 to 18 claim 2 characterised in that the biologically active agent or agents is selected from the group consisting of a drug, peptide, protein, hormone, growth factor, enzyme, polysaccharide, living or dead cells or viruses or parts thereof, plasmids, polynucleotides, water soluble ions, salts and any combination thereof.
- 20. (Currently amended) A bioresorbable sol-gel derived SiO<sub>2</sub>, obtainable according to the method of any of claims 2 to 19 claim 2, characterised in that

- a) the  $SiO_2$  is a monolith, preferably with a minimum diameter of  $\geq 0.5$  mm,
- b) the  $SiO_2$  comprises no biologically active agent other than the  $SiO_2$  itself, and
- c) the dissolution rate of the  $SiO_2$  in a TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.04 wt-%/h, preferably  $\geq$  0.07 wt-%/h and more preferably  $\geq$  0.15 wt-%/h.
- 21. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim
- 2, characterised in that
- a) the  $SiO_2$  is a monolith, preferably with a minimum diameter of  $\geq 0.5$  mm,
- b) the  $SiO_2$  comprises at least one biologically active agent other than the  $SiO_2$  itself, and
- c) the dissolution rate of the  $SiO_2$  in a TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\ge$  0.35 wt-%/h.
- 22. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2, characterised in that

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- a) the  $SiO_2$  is a coating, preferably with a thickness of < 0.5 mm,
- b) the  $SiO_2$  comprises no biologically active agent other than the  $SiO_2$  itself, and
- the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.04 wt-%/h, preferably  $\geq$  0.07 wt-%/h and more preferably  $\geq$  0.15 wt-%/h.
- 23. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2, characterised in that
- a) the  $SiO_2$  is a coating, preferably with a thickness of < 0.5 mm,
- b) the  $\mathrm{SiO}_2$  comprises at least one biologically active agent other than the  $\mathrm{SiO}_2$  itself, and
- the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.04 wt-%/h, preferably  $\geq$  0.07 wt-%/h and more preferably  $\geq$  0.15 wt-%/h.

- 24. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2 characterised in that
- a) the  $SiO_2$  is a particle, preferably with a maximum diameter of  $\leq$  100  $\mu m$ ,
- b) the  $SiO_2$  comprises no biologically active agent other than the  $SiO_2$  itself, and
- c) the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.04 wt-%/h, preferably  $\geq$  0.07 wt-%/h and more preferably  $\geq$  0.15 wt-%/h.
- 25. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2 characterised in that
- a) the  $SiO_2$  is a particle, preferably with a maximum diameter of  $$\leq$~100~\mu m,$
- b) the  $SiO_2$  comprises at least one biologically active agent other than the  $SiO_2$  itself, and
- c) the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.5 wt-%/h.

- 26. (Currently amended) The  $SiO_2$  according to any of claims 20,  $\frac{22}{23}$  and  $\frac{24}{24}$  claim 20, characterised in that the dissolution rate of the  $SiO_2$  is  $\frac{2}{20}$  0.30 wt-%/h.
- 27. (Currently amended) The  $SiO_2$  according to claim 21 or 26, characterised in that the dissolution rate of the  $SiO_2$  is  $\ge 0.5$  wt-%/h preferably  $\ge 1.0$  wt-%/h, more preferably  $\ge 2.0$  wt-%/h and most preferably  $\ge 4.0$  wt-%/h.
- 28. (Currently amended) A bioresorbable sol-gel  $\frac{\text{derivedSiO}_2}{\text{derived SiO}_2}$  obtainable according to the method of  $\frac{\text{any of claims 2}}{\text{to 19}}$  claim 2, characterised in that
- a) the  $SiO_2$  is a monolith, preferably with a minimum diameter of  $\ge 0.5$  mm,
- b) the  $SiO_2$  comprises no biologically active agent other than the  $SiO_2$  itself, and
- the dissolution rate of the SiO<sub>2</sub> in a TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.15 wt-%/h, preferably from 0.002 to 0.07 wt-%/h, and more preferably from 0.006 to 0.05 wt-%/h.

- 29. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim
- 2, **characterised** in that
- a) the  $SiO_2$  is a monolith, preferably with a minimum diameter of  $\geq$  0.5 mm,
- b) the  $SiO_2$  comprises at least one biologically active agent other than the  $SiO_2$  itself, and
- the dissolution rate of the  $SiO_2$  in a TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.06 wt-%/h, preferably from 0.002 to 0.05 wt-%/h, and from 0.006 to 0.025 wt-%/h.
- 30. (Currently amended) The  $SiO_2$  according to claim 22 or 23 characterised in that the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.15 wt-%/h, preferably from 0.002 to 0.07 wt-%/h, and more preferably from 0.006 to 0.05 wt-%/h.
- 31. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2 characterised in that

- a) the  $SiO_2$  is a particle, preferably with a maximum diameter of  $\leq$  100  $\mu m$ ,
- b) the  $SiO_2$  comprises no biologically active agent other than the  $SiO_2$  itself, and
- c) the dissolution rate of the  $SiO_2$  in TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.008, and preferably from 0.002 to 0.003 wt-%/h.
- 32. (Currently amended) A bioresorbable sol-gel derived  $SiO_2$ , obtainable according to the method of any of claims 2 to 19 claim 2 characterised in that
- a) the  $SiO_2$  is a particle, preferably with a maximum diameter of  $\leq$  100  $\mu m$ ,
- b) the  $SiO_2$  comprises at least one biologically active agent other than the  $SiO_2$  itself, and
- the dissolution rate of the SiO<sub>2</sub> in TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.10 wt-%/h, preferably from 0.002 to 0.07 wt-%/h, and more preferably from 0.006 to 0.05 wt-%/h.

- 33. (Currently amended) A bioresorbable sol-gel derived SiO<sub>2</sub> monolith, preferably with a minimum diameter of  $\geq$  0.5 mm, coating, preferably with a thickness of < 0.5 mm, or particle, preferably with a maximum diameter of  $\leq$  100 µm, obtainable according to the method of any of claims 2 to 19 claim 2, wherein said SiO<sub>2</sub> comprises a biologically active agent other than the SiO<sub>2</sub> itself and said biologically active agent is a peptide, protein or cell, characterised in that the dissolution rate of the SiO<sub>2</sub> in TRIS buffer at a temperature of +37 °C and pH 7.4 is  $\geq$  0.04 wt-%/h, preferably  $\geq$  0.07 wt-%/h and more preferably  $\geq$  0.15 wt-%/h.
- 34. (Currently amended) A bioresorbable sol-gel derived SiO<sub>2</sub> monolith, preferably with a minimum diameter of  $\geq$  0.5 mm, coating, preferably with a thickness of < 0.5 mm, or particle, preferably with a maximum diameter of  $\leq$  100 µm, obtainable according to the method of any of claims 2 to 19 claim 2, wherein said SiO<sub>2</sub> comprises a biologically active agent other than the SiO<sub>2</sub> itself and said biologically active agent is a peptide, protein or cell, characterised in that the dissolution rate of the SiO<sub>2</sub> is  $\geq$  0.5 wt-%/h and preferably  $\geq$  4.0 wt-%/h.

- 35. (Currently amended) A bioresorbable sol-gel derived SiO<sub>2</sub> monolith, preferably with a minimum diameter of ≥ 0.5 mm, coating, preferably with a thickness of < 0.5 mm, or particle, preferably with a maximum diameter of ≤ 100 μm, obtainable according to the method of any of claims 2 to 19 claim 2, wherein said SiO<sub>2</sub> comprises a biologically active agent other than the SiO<sub>2</sub> itself and said biologically active agent is a peptide, protein or cell, characterised in that the dissolution rate of the SiO<sub>2</sub> in TRIS buffer at a temperature of +37 °C and pH 7.4 is from 0.001 to 0.15 wt-%/h, preferably from 0.002 to 0.07 wt-%/h, and more preferably from 0.006 to 0.05 wt-%/h.
- 36. (Currently amended) Use of a bioresorbable sol-gel derived SiO<sub>2</sub> according to any of claims 20 to 35 claim 20 for administering a biologically active agent to a human or animal body, wherein said use comprises administering selected from the group consisting of oral, buccal, rectal, parenteral, pulmonary, nasal, ocular, intrauterine, vaginal, urethral, topical, transdermal and surgically implantable administering.

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37. (Currently amended) Use of a bioresorbable sol-gel derived  $SiO_2$  according to any of claims 20 to 35 claim 20 for administering a biologically active agent to a plant.